

### **AMENDMENTS TO THE CLAIMS**

**This listing of claims will replace all prior versions and listings of claims in the application:**

#### **LISTING OF CLAIMS:**

1. (original): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the step of precipitating and isolating parahydroxybenzoic acid in an aqueous solvent at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.
2. (currently amended): The process for preparing crystalline parahydroxybenzoic acid anhydride according to claim 1, wherein the precipitating and isolating step is performed at a temperature which is in the range from the transition temperature to ~~said~~ the transition temperature + 30°C.
3. (original): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the step of precipitating and isolating parahydroxybenzoic acid with acid from a solution of parahydroxybenzoate in an aqueous solvent at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.
4. (original): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the steps of: precipitating parahydroxybenzoic acid in an aqueous solvent with acid, heating the parahydroxybenzoic acid precipitates to dissolve the same, and re-precipitating and isolating the parahydroxybenzoic acid at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.

5. (currently amended): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the steps of:

providing a suspension of parahydroxybenzoic acid monohydrate in an aqueous solvent;  
preparing a solution of parahydroxybenzoic acid in an aqueous solvent by heating the suspension; ~~and~~  
precipitating crystalline parahydroxybenzoic acid anhydride by keeping said solution at a temperature equal to or above the transition temperature of parahydroxybenzoic acid; and  
isolating the crystalline parahydroxybenzoic acid anhydride at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.

6. (currently amended): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the steps of:

~~providing preparing~~ a suspension of parahydroxybenzoic acid monohydrate in an aqueous solvent,  
changing parahydroxybenzoic acid monohydrate to parahydroxybenzoic acid anhydride  
by heating the suspension to a temperature equal to or above the transition temperature of  
parahydroxybenzoic acid, and  
isolating the crystalline parahydroxybenzoic acid anhydride at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.

7. (currently amended): The process for preparing crystalline parahydroxybenzoic acid anhydride according to ~~any one of claims 1 to 6~~ claim 1, 2, 3, 4, 5 or 6, wherein the aqueous solvent is water and the transition temperature of parahydroxybenzoic acid is 52 to 54°C.

8. (currently amended): Crystalline parahydroxybenzoic acid anhydride, wherein the specific surface area of particles that can pass through a 100 mesh (150  $\mu\text{m}$ ) sieve and can not pass through a 140 mesh (106  $\mu\text{m}$ ) sieve is equal to or less than 0.3  $\text{m}^2/\text{g}$ .

9. (original): The crystalline parahydroxybenzoic acid anhydride according to claim 8, wherein the angle of repose is equal to or less than 45°.

10. (original): The crystalline parahydroxybenzoic acid anhydride according to claim 8 or 9, wherein the compression ratio calculated according to the following formula is equal to or less than 10%:  $(\text{packed bulk density} - \text{aerated bulk density}) / \text{packed bulk density} \times 100$ .